

K112283

OCT 20 2011



1430 Waukegan Road  
McGraw Park, IL 60085

[www.cardinal.com](http://www.cardinal.com)

### SMDA REQUIREMENTS

#### **510(k) SUMMARY OF SAFETY AND EFFECTIVENESS** **DuraBlue™ Sterilization Wrap**

Manufacturer: Cardinal Health 200, LLC  
1430 Waukegan Road  
McGraw Park, IL 60085

Regulatory Affairs Contact: Lavenia Ford  
1430 Waukegan Road  
McGraw Park, IL 60085

Telephone Number: (847) 887-3323

Date summary Prepared: September 28, 2011

Trade Name: DuraBlue™ Sterilization Wrap

Classification: Class II per 21 CFR § 880.6850

Classification Name: Sterilization Wrap

Predicate Device: K092167- KIMGUARD ONE-STEP Sterilization Wrap (Models KC100, KC200, KC400, KC500 and KC600) for the Use With the Lumen Cycle of the Amsco V – PRO 1 Low Temperature Sterilization System and With the Lumen and Non Lumen Cycles of the Amsco V – PRO 1 Plus Low Temperature Sterilization System

Description:

Cardinal Health DuraBlue™ Sterilization Wrap is double layer sterilization wraps made from 100% polypropylene spunbond-meltblown-spunbond (SMS) fabric. The wrap is intended to be used to enclose another medical device that is to be sterilized by a health care provider by STERRAD® 100S System. This wrap design allows for use of the simultaneous double-wrapping technique and also allows for a sterilized pack to be opened aseptically.

This submission covers six different models of Cardinal Health DuraBlue™ Sterilization Wrap. Each model is made from material of a different basis weight, though all models utilize the same material technology.

Extensive performance testing has been completed on Cardinal Health DuraBlue™ Sterilization Wrap. Successful completion of the sterilization performance tests listed below demonstrated that the wrap both allows for sterilization of the enclosed contents and maintains sterility of the enclosed contents for at least 30 days.

**Indications for Use:**

Cardinal Health DuraBlue™ Sterilization Wrap is intended to be used to enclose another medical device that is to be sterilized by a health care provider by STERRAD® 100S System. The wrap is intended to allow sterilization of the enclosed medical device(s) and to maintain sterility of the enclosed devices for 30 days. All models have been validated for sterilization of lumens 2.5 mm in diameter or larger and 250 mm in length or less.

**Wrap Model Recommendations<sup>1</sup>**

Sterilization Wrap Model	Intended Load	Maximum Recommended Wrapped Package Content Weights <sup>2</sup>
CH100	Very light weight package (for example: batteries).	3 lbs (1.36 kg)
CH200	Light weight package (for example: telescope with light cord).	6 lbs (2.7 kg)
CH300	Light to moderate weight package (for example: general use medical instruments).	9.7 lbs (4.4 kg)
CH400	Moderate to heavy weight package (for example: general use medical instruments).	9.7 lbs (4.4 kg)
CH500	Heavy weight package (for example: general use medical instruments).	9.7 lbs (4.4 kg)
CH600	Very heavy weight package (for example: general use medical instruments).	9.7 lbs (4.4 kg)

- CH100: Metal instruments
- CH200: 15 in. x 10 in. x 1.2 in. tray containing metal instruments
- CH300: 15 in. x 10 in. x 1.2 in. tray containing metal instruments
- CH400: 15 in. x 10 in. x 1.2 in. tray containing metal instruments
- CH500: 15 in. x 10 in. x 1.2 in. tray containing metal instruments
- CH600: 15 in. x 10 in. x 1.2 in. tray containing metal instruments

**Note:** The loads used in the Sterility Maintenance Validation Study corresponded to the maximum wrapped package content weights in Table 2.

<sup>1</sup>Individual results may differ due to factors such as variations in handling practices, wrapping techniques, and folding methods. Results may also differ due to the use of irregularly shaped contents, which may put added stress on the wrap. Each healthcare facility should determine for itself which wrap model is most appropriate for each intended use.

<sup>2</sup> It is recommended to not exceed the maximum wrapped package content weights indicated for each wrap model. Furthermore, it is recommended to not exceed the number, weight and size of individual content types that were validated for the DuraBlue™ Sterilization Wraps.

### Substantial Equivalence

The DuraBlue™ Sterilization Wrap is substantially equivalent to the predicate devices.

- Both devices are double layer sterilization wraps which allow for use of the simultaneous double-wrapping technique and for a sterilized pack to be opened aseptically.
- Both devices are intended to be used with vaporized hydrogen peroxide sterilizers that operate at low temperatures and pressures.
- Both devices are available in six comparable models of varying basis weights, which are recommended for use with the same maximum content weights.
- Both devices have the same dimensional specifications.
- Both devices are 100% polypropylene spunbond-meltblown-spunbond (SMS) trilaminate nonwoven fabric.
- Both devices demonstrate maintenance of package sterility for at least 30 days following sterilization by STERRAD® 100S.
- Performance and safety attributes are substantially equivalent to the predicate. The physical properties of all wrap models have been characterized both before and after exposure to STERRAD® 100S sterilization. The resulting data supports the conclusion that Cardinal Health DuraBlue™ Sterilization Wrap is substantially equivalent to the predicate, and the DuraBlue™ Sterilization Wraps are compatible with the identified STERRAD®100S System.

### Summary of Testing

DuraBlue™ Sterilization Wrap performance has been tested in accordance with the applicable requirements recommended in the FDA's Guidance Document [*Premarket Notification 510(k) Submissions for Medical Sterilization Packaging System in Health Care Facilities; Draft Guidance for Industry and FDA*] (March 7, 2002). Testing included sterilization efficacy, event related maintenance of package sterility, physical properties, and biocompatibility in compliance with the methods of ISO 10993. Data from testing demonstrates that the performance of the DuraBlue™ Sterilization Wrap is substantially equivalent to that of Kimberly-Clark KIMGUARD ONE-STEP Sterilization Wrap.



## DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration  
10903 New Hampshire Avenue  
Document Control Room -WO66-G609  
Silver Spring, MD 20993-0002

Cardinal Health 200, LLC  
% Mr. Ned Devine  
Senior Staff Engineer  
Underwriters Laboratories, Incorporated  
333 Pfingsten Road  
Northbrook, Illinois 60062

OCT 20 2011

Re: K112283

Trade/Device Name: Cardinal Health DuraBlue™ Sterilization Wrap  
Regulation Number: 21 CFR 880.6850  
Regulation Name: Sterilization Wrap  
Regulatory Class: II  
Product Code: FRG  
Dated: October 4, 2011  
Received: October 5, 2011

Dear Mr. Devine:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

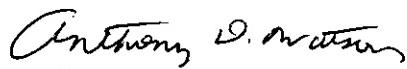
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please go to <http://www.fda.gov/AboutFDA/CentersOffices/CDRH/CDRHOffices/ucm115809.htm> for the Center for Devices and Radiological Health's (CDRH's) Office of Compliance. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address <http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,

  
Anthony D. Watson, B.S., M.S., M.B.A.  
Director  
Division of Anesthesiology, General Hospital,  
Infection Control and Dental Devices  
Office of Device Evaluation  
Center for Devices and Radiological Health

Enclosure

## Indications for Use

510(k) Number (if known): K112283

Device Name: Cardinal Health DuraBlue™ Sterilization Wrap

STERRAD® 100S

Indications for Use:

Cardinal Health DuraBlue™ Sterilization Wrap is intended to be used to enclose another medical device that is to be sterilized by a health care provider by the STERRAD® 100S System. The wrap is intended to allow sterilization of the enclosed medical device(s) and to maintain sterility of the enclosed devices for 30 days. All models have been validated for sterilization of lumens 2.5 mm in diameter or larger and 250 mm in length or less.

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Prescription Use \_\_\_\_\_ AND/OR Over-The-Counter Use X \_\_\_\_\_  
(Part 21 CFR 801 Subpart D) (21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

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Concurrence of CDRH, Office of Device Evaluation (ODE)

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(Division Sign-Off)

Division of Anesthesiology, General Hospital  
Infection Control, Dental Devices

510(k) Number: K112283